AMENDED CLAIM SET:

1. (currently amended) An optically active salicylideneaminoalcohol compound of formula (1):

$$X_1$$
 X_2
 X_1
 X_2
 X_1
 X_2
 X_1
 X_2
 X_3
 X_4
 X_4
 X_4
 X_5
 X_6
 X_6
 X_7
 X_8
 X_8
 X_8
 X_8
 X_8
 X_8
 X_8
 X_8
 X_8
 X_8

wherein

R₁ represents an alkyl group which may be substituted with a group selected from an alkoxy group, an aralkyloxy group, an aryloxy group and cycloalkoxy group, an aralkyl, aryl or cycloalkyl group all of which may be substituted with a group selected from an alkyl group, an alkoxy group, an aralkyloxy group, an aryloxy group roup and a cycloalkoxy group,

R₂ represents an alkyl group, a cycloalkyl group, or an aralkyl or phenyl group which may be substituted with a group selected from an alkyl group, an alkoxy group, an aralkyloxy group, an aryloxy group and a cycloalkoxy group,

when X_1 represents a nitro group, X_2 is a hydrogen atom, when X_1 represents a chlorine atom, X_2 is a chlorine atom, and when X_1 is a hydrogen atom, X_2 is a fluorine atom; and

the carbon atom denoted by " * " is an asymmetric carbon atom having either an S or R configuration.

2. (original) An optically active salicylideneaminoalcohol compound according to claim 1, wherein R_1 and R_2 are the same or different and independently represent an alkyl group, an aralkyl group, a phenyl group, a 2-methoxyphenyl group, a 2-tert-butylphenyl group or a 2-octyloxy-5-tert-butylphenyl group.

3. (currently amended) A process for producing an optically active salicylideneaminoalcohol compound as defined in claim 1, which comprises

reacting an optically active amino alcohol of formula (2):

$$R_1$$
 R_2 R_2

wherein R₁ represents an alkyl group which may be substituted with a group selected from an alkoxy group, an aralkyloxy group, an aryloxy group and cycloalkoxy group, an aralkyl, aryl or cycloalkyl group all of which may be substituted with a group selected from an alkyl group, an alkoxy group, an aralkyloxy group, an aryloxy group roup, and a cycloalkoxy group, R₂ represents a hydrogen atom, an alkyl group, a cycloalkyl group or an aralkyl or phenyl group which may be substituted with a group selected from an alkyl group, an alkoxy group, an aralkyloxy group, an aryloxy group roup and a cycloalkoxy group, and the carbon atom denoted by " * " is an asymmetric carbon atom having either an S or R configuration,

3

with a 2-hydroxybenzaldehyde derivative of formula (3):

$$X_1$$
—CHO
$$X_2$$

$$X_3$$

$$X_3$$

$$X_4$$

$$X_4$$

$$X_5$$

$$X_6$$

$$X_1$$

wherein when X_1 represents a nitro, X_2 is a hydrogen atom, when X_1 represents a chlorine atom, X_2 is a chlorine atom, and when X_1 is a hydrogen atom, X_2 is a fluorine atom.

- 4. (original) A process according to claim 3, wherein R_1 and R_2 are the same or different and independently represent an alkyl group, an aralkyl group, a phenyl group, a 2-methoxyphenyl group, a 2-tert-butoxy-5-tert-butylphenyl group or a 2-octyloxy-5-tert-butylphenyl group.
- 5. (original) A chiral copper complex obtained by contacting a mono-valent or di-valent copper compound with an optically active salicylideneaminoalcohol compound as defined in claim 1 or 2.
- 6. (previously presented) An adduct comprising a chiral copper complex as defined in claim 5 and a prochiral olefin of formula (5):

$$\begin{array}{c|c}
R_3 & R_5 \\
R_4 & R_6
\end{array}$$
(5)

wherein R₃, R₄, R₅ and R₆ independently represent a hydrogen atom, a halogen atom, a (Cl-C10)alkyl group which may be substituted with a halogen atom or a lower alkoxy group, a (C4-C8)cycloalkyl group, an aryl group which may be substituted with a halogen atom or a lower

4

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alkoxy group, or an alkoxy group; or R_3 and R_4 , or R_5 and R_6 together form a cycloalkylene group having 2-4 carbon atoms, provided that one of R_3 , R_4 , R_5 and R_6 groups represents an alkenyl group which may be substituted with a halogen atom, an alkoxy group or an alkoxy carbonyl group, of which alkoxy may be substituted with a halogen atom or atoms, and provided that when R_3 and R_5 are the same, R_4 and R_6 are not the same.

7. (original) A method for producing a chiral copper complex of formula (1):

$$X_1$$
 X_2
 R_2
 R_2

wherein R_1 and R_2 are the same or different and independently represent an alkyl group, an aralkyl group, a phenyl group, a 2-methoxyphenyl group, a 2-tert-butoxy-5-tert-butylphenyl group, or a 2-octyloxy-5-tert-butylphenyl group, when X_1 represents a nitro group, X_2 is a hydrogen atom, when X_1 represents a chlorine atom, X_2 is a chlorine atom, and when X_1 represents a hydrogen atom, X_2 is a fluorine atom, the carbon atom denoted by " * " is an asymmetric carbon atom having either an S or R configuration,

which comprises contacting a di-valent copper compound, in an inert organic solvent, with a chiral salicylideneaminoalcohol compound of formula (1):

Application No.: 10/625,604

Docket No.: 2185-0703P

$$X_1$$
 X_2
 R_1
 R_2
 R_2
 R_2
 R_2
 R_2
 R_3
 R_4
 R_2
 R_2
 R_3
 R_4
 R_2
 R_3
 R_4
 R_5
 R_5
 R_7
 R_7
 R_8

wherein R_1 , R_2 X_1 , X_2 and " * " respectively have the same meaning as defined above.

- 8. (original) A method according to claim 7, which further comprises subjecting the resulting solution to precipitation and collecting the precipitated crystals of said chiral copper complex of formula (1)'.
- 9. (original) A method according to claim 8, said precipitation is carried out by cooling the reaction solution or by adding an aliphatic hydrocarbon solvent.
 - 10. (cancelled).
 - 11. (cancelled).
 - 12. (cancelled).